

UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address COMMISSIONER FOR PATENTS FO Box 1430 Alexandria, Virginia 22313-1450 www.tepto.gov

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/081,953	02/22/2002	William J. Hennen	2820-4428.2US	6427	
95261 Durham, Jones	7590 08/17/201 & Pinegar	EXAMINER			
Intellectual Pro	perty Law Group	CHEN, STACY BROWN			
P.O. Box 4050 Salt Lake City			ART UNIT	PAPER NUMBER	
			1648		
			NOTIFICATION DATE	DELIVERY MODE	
			08/17/2010	ELECTRONIC	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Notice of the Office communication was sent electronically on above-indicated "Notification Date" to the following e-mail address(es):

patents@djplaw.com kolsen@djplaw.com cwickstrand@djplaw.com

Advisory Action Before the Filing of an Appeal Brief

Application No.	Applicant(s)		
10/081,953	HENNEN ET AL.		
Examiner	Art Unit		
01 0 01	4040		
Stacy B. Chen	1648		

	Stacy B. Chen	1040	
The MAILING DATE of this communication appe	ars on the cover sheet with the o	orrespondence add	ress
THE REPLY FILED 02 August 2010 FAILS TO PLACE THIS AF	PLICATION IN CONDITION FOR	ALLOWANCE.	
 X The reply was filed after a final rejection, but prior to or on application, applicant must timely file one of the following application in condition for allowance; (2) a Notice of Appe for Continued Examination (RCE) in compliance with 37 C periods: 	replies: (1) an amendment, affidavi	t, or other evidence, w with 37 CFR 41.31; or	hich places the (3) a Request
 a) The period for reply expiresmonths from the mailing 			
 The period for reply expires on: (1) the mailing date of this A no event, however, will the statutory period for reply expire to 	iter than SIX MONTHS from the mailing	date of the final rejection	n.
Examiner Note: If box 1 is checked, check either box (a) or (MONTHS OF THE FINAL REJECTION. See MPEP 706.07(I).		
Extensions of time may be obtained under 37 CFR 1.136(a). The date have been filled is the date for purposes of determining the period of ext under 37 CFR 1.17(a) is calculated from: (1) the expiration date of the set set forth in (b) above, if checked. Any reply received by the Office later may reduce any earned patient term adjustment. See 37 CFR 1.704(b). NOTICE OF APPEAL.	ension and the corresponding amount of hortened statutory period for reply origing than three months after the mailing date	of the fee. The appropria nally set in the final Office	ate extension fee e action; or (2) as
2. The Notice of Appeal was filed on A brief in comp	liance with 37 CFR 41.37 must be t	iled within two months	s of the date of
filing the Notice of Appeal (37 CFR 41.37(a)), or any exter Notice of Appeal has been filed, any reply must be filed wi	sion thereof (37 CFR 41.37(e)), to	avoid dismissal of the	appeal. Since
<u>AMENDMENTS</u>			
3. The proposed amendment(s) filed after a final rejection, b			cause
(a) They raise new issues that would require further cor		E below);	
(b) They raise the issue of new matter (see NOTE below			
 (c) They are not deemed to place the application in beti appeal; and/or 	ter form for appeal by materially rec	lucing or simplifying ti	ie issues ioi
(d) ☐ They present additional claims without canceling a c	corresponding number of finally reje	cted claims.	
NOTE: (See 37 CFR 1.116 and 41.33(a)).	,,		
4. The amendments are not in compliance with 37 CFR 1.12	1. See attached Notice of Non-Co.	mpliant Amendment (I	PTOL-324)
5. Applicant's reply has overcome the following rejection(s):		- ipinani i i i i i i i i i i i i i i i i i	
Newly proposed or amended claim(s) would be all non-allowable claim(s).		imely filed amendmer	nt canceling the
 For purposes of appeal, the proposed amendment(s): a) [how the new or amended claims would be rejected is proving the proposed amendment of the proposed		be entered and an e	xplanation of
The status of the claim(s) is (or will be) as follows:			
Claim(s) allowed:			
Claim(s) objected to: Claim(s) rejected:			
Claim(s) rejected: Claim(s) withdrawn from consideration:			
AFFIDAVIT OR OTHER EVIDENCE			
The affidavit or other evidence filed after a final action, but because applicant failed to provide a showing of good and			
was not earlier presented. See 37 CFR 1.116(e).			
 The affidavit or other evidence filed after the date of filing entered because the affidavit or other evidence failed to o showing a good and sufficient reasons why it is necessary 	vercome <u>all</u> rejections under appea	l and/or appellant fail:	s to provide a
10. The affidavit or other evidence is entered. An explanation	of the status of the claims after er	ntry is below or attach	ed.
REQUEST FOR RECONSIDERATION/OTHER		•	
 The request for reconsideration has been considered but See Continuation Sheet. 	does NOT place the application in	condition for allowan	ce because:
12. Note the attached Information Disclosure Statement(s). (PTO/SB/08) Paper No(s)		
13. Other:			
	(Story B. Chan)		
	/Stacy B Chen/ Primary Examiner, Art U	nit 16/18	
	Filliary Examiner, Art U	THE 1040	

Continuation of Item 11

Claims 1-16 and 18-22 remain rejected under 35 U.S.C. 102(e) as being anticipated by Dopson (PGPub 2002/0044942A1, "Dopson", published April 18, 2002, with priority to provisional application 60/233,400, filed September 18, 2000), for reasons of record. Applicant indicates in the response filed August 2, 2010, that an affidavit may be filed once all other issues in this application are resolved.

Claims 1-16 and 18-22 are rejected under 35 U.S.C. 103(a) as being unpatentable over Tokoro (US Patent 5,080,895) in view of Kirkpatrick et al. (US Patent 5,840,700, "Kirkpatrick"), for reasons of record. Applicant's arguments have been carefully considered but fail to persuade. Applicant's arguments are directed to the following.

Applicant points out that Tokoro admits that "the immunological functions of the transfer factor-like component...are not known", see Tokoro, col. 7, lines 44-47. Applicant argues that in view of the lack of knowledge of the immunological function of Tokoro's transfer factor-like component, one would not have been able to predict that it would be useful to induce a T-cell mediated immune response.

In response to Applicant's argument, the Office agrees that Tokoro did not know the exact immunological functions of their transfer factor-like fraction. However, that statement by Tokoro is simply saying that the particulars of how the transfer factor-like component inspected the immune system were not known, not that it was not useful to a subject's immune system. In col. 7, lines 30-43, the transfer factor-like component is clearly disclosed as useful for teatment of diseases and additives in food.

Applicant maintains the position that the predictability of whether an antigen would induce a T-cell response was low.

In response to Applicant's argument, the Office recognizes that not every antigen will induce a T-cell response, such as certain bacterial antigens suggested by Tokoro that do not induce T-cell mediated immunity in hens. The Office also acknowleds that a T-cell response is critical to the production of transfer factor. However, Tokoro does not have to teach that the antigens elicit a T-cell response. The obviousness rejection relies not Tokoro's description of a transfer factor-like component that is specific for an aptropen, in combination with a reference that teaches an antigen/pathogen that is known to induce T-cell mediated immunity in an animal. For example, Tokoro falls to disclose EBV-specific transfer factor-like component, including those other than intestinal infectious diseases (col. 4, lines 16-18). Antigens from pollen, bacteria, virusee, molds, allergens, blood from affected animals, sperm and toknown may be used in the production of transfer factor-like component (col. 4, lines 35-57). One would have been motivated to select an antigen from a clinically significant pathogen such as Epstein-Barr virus, a known pathogen for which a vaccine is desirable to prevent mononucleosis (Kirtipatrick, col. 5, lines 7-30). By immunizing hens with an EBV antigen, Tokoro's hens would have produced transfer factor-like component specific for EBV.